



## VIRTUAL REALITY IN PHARMACY EDUCATION: A REVIEW OF APPLICATIONS, ADVANTAGES, AND FUTURE PROSPECTS

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### ABSTRACT

Pharmacy education relies heavily on practical demonstrations to build student understanding of drug actions, physiological responses, and experimental techniques. Traditionally, this has depended on live animal models – an approach increasingly challenged by ethical concerns, regulatory restrictions under the 3Rs principle (Replacement, Reduction, Refinement), high infrastructure costs, and inconsistent resource availability across institutions. Virtual Reality (VR) technology has emerged as a transformative educational tool capable of simulating immersive laboratory environments and demonstrating complex pharmaceutical concepts with unprecedented depth and specificity. This comprehensive review examines the published literature on the application of VR and simulation-based learning across five key pharmaceutical streams: (i) Pharmacology – drug-receptor binding, ion channel modulation, dose-response relationships; (ii) Pharmaceutics – dosage form manufacturing, GMP environments, dissolution and release mechanisms; (iii) Pharmaceutical Chemistry – molecular-level reaction visualisation, synthesis, and spectroscopic analysis; (iv) Pharmaceutical Biotechnology – genetic engineering, recombinant protein production, cell culture, immunology; and (v) Clinical Pharmacy – patient counselling, prescription error detection, and medication management. Evidence from multiple studies confirms that VR enhances student engagement, improves knowledge retention, supports procedural understanding, and provides equitable access to quality practical learning. The review further discusses limitations of current VR tools and the future potential of immersive technologies as mainstream supplements to conventional pharmacy practical training.

**KEYWORDS:** Virtual reality, Pharmacy education, Pharmacology, Pharmaceutics, Pharmaceutical chemistry, Biotechnology, Clinical pharmacy, Drug-receptor interaction, Molecular visualisation, Immersive learning, 3Rs principle, Genetic engineering, Dosage form simulation.

### INTRODUCTION

#### Pharmacy Education and the Role of Practical Learning

Pharmacy education is a discipline that demands both rigorous theoretical knowledge and competent practical skills. The study of pharmacology – the branch of pharmaceutical sciences concerned with drug mechanisms, pharmacokinetics, pharmacodynamics, and dose-response relationships – sits at the core of

pharmacy training. A thorough understanding of pharmacology cannot be achieved through classroom instruction alone; practical exposure to experimental procedures is essential for students to observe and internalize drug effects on living system.<sup>[1]</sup>

Conventionally, pharmacology practical sessions in Indian pharmacy colleges involve the use of animal models such as rats, mice, rabbits, and frogs to

demonstrate drug actions. These experiments provide first-hand observation of phenomena such as changes in heart rate and blood pressure, smooth muscle contraction and relaxation, central nervous system depression or stimulation, and dose-response relationships. Such direct observation is irreplaceable in building the intuitive, experiential understanding that distinguishes a pharmacology-literate practitioner from one who has only read about drug actions.<sup>[2]</sup>

### **Limitations of Conventional Pharmacology Practical Training**

Despite its educational value, animal-based pharmacology practical training faces compounding challenges in modern educational settings. The infrastructure required – dedicated animal houses, physiological recording equipment, ethical clearance from Institutional Animal Ethics Committees (IAEC), trained veterinary staff, and ongoing animal care costs – places a significant financial burden on institutions.<sup>[3]</sup> Many pharmacy colleges, particularly those in smaller cities and rural Maharashtra, operate with limited laboratory budgets and are unable to sustain a fully equipped pharmacology practical programme.

Equally important are the ethical dimensions of animal use in education. The 3Rs framework – Replacement, Reduction, and Refinement – originally proposed by Russell and Burch in 1959, has been adopted globally as the ethical standard for minimising animal use in experimental science. In India, the Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA) has progressively strengthened regulations governing animal use in educational institutions. As a result, numerous colleges have curtailed or suspended animal-based practicals, creating a gap in practical pharmacology training that students increasingly fill only through theoretical study.<sup>[4]</sup>

Beyond infrastructure and ethics, traditional practical sessions suffer from pedagogical limitations. Animal responses are inherently variable; time constraints restrict the number of experiments students can observe; and the passive, observational role most students occupy during group practicals limits active learning engagement. These limitations collectively argue for the development and adoption of credible alternative practical learning tools.<sup>[5]</sup>

### **Virtual Reality as an Emerging Educational Technology**

Virtual Reality (VR) is a computer-generated technology that creates immersive three-dimensional environments in which users can interact with simulated objects and scenarios in real time. Since its early applications in aviation and military training simulation, VR has been increasingly adopted across education, healthcare, and industry. In the health sciences, VR has been used to simulate surgical procedures, clinical patient encounters, anatomical dissection, and laboratory experiments.<sup>[6]</sup>

For pharmacy education specifically, VR offers a unique combination of advantages: the ability to demonstrate pharmacological experiments without animals; the capacity for unlimited repetition without resource expenditure; visual representation of microscopic drug-receptor interactions and physiological processes that cannot be seen in a conventional laboratory; and accessibility via devices ranging from dedicated VR headsets to standard smartphones.<sup>[7]</sup> This review surveys the evidence on how VR has been applied in pharmacy and health sciences education, what outcomes it has achieved, and what its future role in pharmacology practical training may be.

### **Overview of Virtual Reality Technology in Education Types of VR Systems Used in Education**

VR systems used in educational settings range from high-immersion head-mounted displays (HMDs) such as the Meta Quest and HTC Vive, which deliver stereoscopic 3D visuals and motion tracking, to desktop-based virtual environments accessible through standard computers, to mobile VR experiences using smartphones and low-cost cardboard or plastic headsets. Each category offers a different balance of immersion, cost, and accessibility.<sup>[6]</sup>

High-immersion HMD systems provide the most realistic virtual experience, with six degrees of freedom (6DoF) movement tracking allowing students to physically lean, reach, and interact within the virtual environment. Studies have consistently linked higher immersion to greater engagement and stronger transfer of learning to real-world tasks.<sup>[8]</sup> However, the cost of HMD hardware – typically INR 25,000–100,000 per unit – limits their adoption in resource-constrained institutions. Desktop and mobile VR platforms sacrifice some immersion but are accessible at minimal or no additional hardware cost, making them more practical for large-scale pharmacy education deployment.<sup>[9]</sup>

### **Augmented Reality vs. Virtual Reality in Pharmacy Education**

Augmented Reality (AR) overlays digital information onto the real-world environment, typically through a smartphone camera or smart glasses. Unlike fully immersive VR, AR preserves the user's physical surroundings while adding interactive digital elements. In pharmacy education, AR has been used to overlay 3D molecular structures onto physical drug samples, annotate laboratory equipment with contextual information, and visualise drug absorption pathways in anatomical context. Studies comparing AR and VR in health science education suggest that both improve learning outcomes relative to conventional instruction, with VR demonstrating advantages in procedural learning and AR excelling in contextual information overlay.<sup>[7,10]</sup>

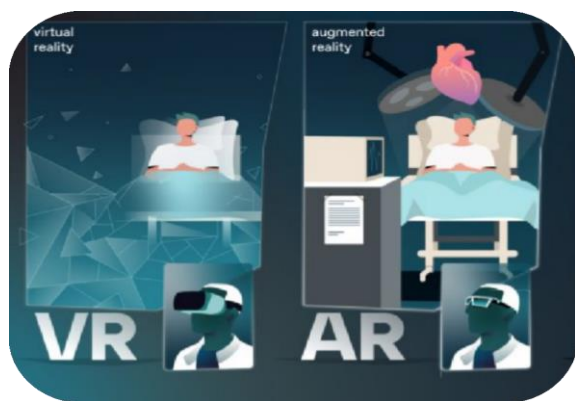


Fig. 1: VR vs AR.

## Applications of VR in Overall Pharmacy Education I: Pharmacology

### 1 Introduction to VR in Pharmacology

Pharmacology is the foundational science of pharmacy – the study of how drugs interact with biological systems to produce their therapeutic or toxic effects. Pharmacology practical training traditionally requires animal models to demonstrate dose-response relationships, autonomic drug effects, cardiovascular pharmacology, CNS drug actions, and smooth muscle pharmacology. The progressive tightening of CPCSEA guidelines in India has significantly curtailed animal use in educational pharmacology, creating an urgent need for validated alternative practical tools.<sup>[3]</sup>

What makes VR uniquely suited to pharmacology is its ability not only to simulate the observable outcomes of drug administration – changes in heart rate, blood pressure, or muscle tone – but also to make the underlying molecular mechanisms visible. For the first time in the history of pharmacy education, students can watch a drug molecule approach a receptor, bind to its active site, trigger a conformational change, activate a G-protein, stimulate adenylyl cyclase, and observe the rise in cyclic AMP concentration that ultimately produces the macroscopic physiological response they are studying. This dual-level visualisation – from molecule to organism – transforms pharmacology from a discipline of observing outcomes to one of truly understanding mechanisms.<sup>[11]</sup>

### 2 Specific VR Applications in Pharmacology

#### a) Drug-Receptor Binding Simulation

VR platforms simulate the entire sequence of receptor pharmacology at atomic resolution. In a virtual pharmacology laboratory, students can observe a rendered three-dimensional model of a G-protein coupled receptor (GPCR) – such as the beta-1 adrenoceptor in cardiac muscle – embedded in the phospholipid bilayer membrane. The student selects a drug such as adrenaline (epinephrine) and watches as the rendered drug molecule approaches the receptor through the extracellular space, enters the orthosteric binding pocket, and undergoes induced-fit binding. The simulation visualises hydrogen bonds forming between

the drug's hydroxyl groups and amino acid residues in the receptor binding site, as well as hydrophobic interactions between the drug's aromatic ring and non-polar pocket residues.<sup>[11]</sup>

Following binding, students observe the conformational change in the receptor's transmembrane helices that activates the associated Gs-protein – watching the alpha subunit exchange GDP for GTP, dissociate from the beta-gamma dimer, and stimulate membrane-bound adenylyl cyclase. The resulting intracellular cascade – rising cAMP, activation of protein kinase A, phosphorylation of regulatory proteins – is animated in real time, culminating in the observable physiological effect: increased heart rate and contractility. Platforms such as Virtual Cell, BioDigital Human, and PharmacologyLab VR have implemented these cascades for multiple receptor systems including adrenoceptors, muscarinic receptors, GABA-A ion channels, and opioid receptors.<sup>[3]</sup>

#### b) Ion Channel Pharmacology

VR enables the visualisation of ion channel pharmacology that is entirely impossible to observe in a conventional laboratory. Students can examine a three-dimensional rendering of a voltage-gated sodium channel in a neuronal membrane and watch local anaesthetic drugs such as lidocaine access the channel through the inner mouth, bind within the pore, and physically block sodium ion flow. The simulation demonstrates the use-dependent nature of this block by showing how repeated channel opening – simulated by applying a train of membrane depolarisations – increases lidocaine binding probability, explaining the clinical phenomenon of frequency-dependent block.<sup>[13]</sup>

Similarly, GABA-A receptor pharmacology – the mechanism of benzodiazepines, barbiturates, and general anaesthetics – is visualised through animated cross-sections of the pentameric ion channel, showing allosteric modulator binding sites distinct from the GABA binding site, and demonstrating how drug binding increases the frequency or duration of chloride ion channel opening.<sup>[3,15]</sup>

#### c) Dose-Response Experiments

VR pharmacology platforms replicate the classic dose-response experiments of pharmacology practical training. Students select a drug, a virtual tissue preparation (isolated ileum, isolated heart, frog sciatic nerve preparation), and a dosing protocol. The simulation generates dose-response curves in real time as the student administers cumulative log doses, with the virtual preparation displaying appropriate responses: smooth muscle contraction, cardiac rate changes, or action potential generation. Students can then add a competitive antagonist – visualising it competing with the agonist at the receptor binding site – and observe the rightward shift of the dose-response curve, directly experiencing the concept of competitive antagonism as a molecular

competition rather than an abstract equation.<sup>[4]</sup>

#### d) Autonomic Pharmacology and Cardiovascular Simulations

Autonomic pharmacology – the study of drugs acting on the sympathetic and parasympathetic nervous systems – is particularly well served by VR. Students can navigate a virtual representation of the autonomic nervous system, observe preganglionic and postganglionic neurons, and watch neurotransmitter release and receptor activation at both ganglionic and effector organ synapses. The effects of drugs such as atropine (muscarinic antagonist), neostigmine (acetylcholinesterase inhibitor), prazosin (alpha-1 antagonist), and propranolol (beta antagonist) are demonstrated at both the molecular and whole-organ level – showing drug binding at the receptor while simultaneously displaying changes in heart rate, blood pressure, pupil diameter, and gastrointestinal motility.<sup>[4,11]</sup>



**Fig. 2: VR in Pharmacology.**

#### 3 Advantages of VR in Pharmacology

VR makes the pharmacological mechanism simultaneously visible at both the molecular and physiological level, a dual perspective that no conventional laboratory tool can provide. The ability to connect drug-receptor binding at the nanometre scale to heart rate changes observed at the organismal level constitutes a genuinely new educational capacity. Studies confirm that this mechanism-level visualisation produces deeper conceptual understanding of pharmacodynamics than whole-animal observation alone.<sup>[3,8]</sup> VR also eliminates animal use entirely, ensuring full compliance with the 3Rs principle and CPCSEA guidelines, while providing an unlimited number of experimental repetitions at no additional cost.<sup>[5]</sup> Students can explore the entire pharmacological space – varying dose, route, time of administration, and patient parameters – within a single practical session.

#### 4 Limitations of VR in Pharmacology

VR pharmacology platforms cannot replicate the variability inherent in biological systems – the unpredictable responses of real animals that build the critical thinking skills of experimental pharmacologists. The tactile skills of animal handling, intraperitoneal injection technique, and organ bath manipulation require

genuine physical practice. Current platforms also do not uniformly cover all drug classes relevant to the Indian B.Pharm pharmacology curriculum, and assessment frameworks for VR-based pharmacology competency have not yet been established by the Pharmacy Council of India.<sup>[5,15]</sup>

## II: Pharmaceutics

### 1 Introduction to VR in Pharmaceutics

Pharmaceutics is the science of formulating drugs into dosage forms suitable for safe, effective, and patient-acceptable administration. It encompasses preformulation science, formulation development, dosage form design, biopharmaceutics, and pharmaceutical manufacturing technology. Practical training in pharmaceutics requires access to industrial-scale and laboratory-scale manufacturing equipment – tablet compression machines, fluid bed dryers, coating pans, high shear granulators, filling lines, and stability chambers – alongside quality control instruments including dissolution apparatus, hardness testers, HPLC systems, and particle size analysers. The capital cost of equipping a pharmaceutics laboratory to the standard required for meaningful practical training is substantial and is beyond the reach of many Indian pharmacy colleges.<sup>[16]</sup>

Virtual Reality addresses this gap by simulating pharmaceutical manufacturing environments, formulation laboratory procedures, and quality control workflows with fidelity sufficient for meaningful skill development. Critically, VR enables visualisation of the physicochemical processes that underlie dosage form performance – dissolution, drug release, membrane permeation, and bioavailability – at scales ranging from molecular interactions in tablet matrices to macroscopic dissolution profiles. This makes VR uniquely valuable in pharmaceutics, where understanding why a dosage form performs as it does requires insight into processes that are invisible in a conventional laboratory.<sup>[17]</sup>

### 2 Specific VR Applications in Pharmaceutics

#### a) Dosage Form Manufacturing Simulation – Tablets

VR manufacturing simulations take students through the complete tablet manufacturing process from a GMP-compliant perspective. In a virtual pharmaceutics laboratory, students begin by performing virtual preformulation studies – observing the crystal structure of a drug substance, measuring its virtual flowability and compressibility characteristics through simulated bulk density and Carr's Index measurements, and testing its compatibility with common excipients through visualised differential scanning calorimetry (DSC) thermograms. The simulation visualises drug-excipient interactions at the molecular level: hydrogen bond formation between drug and binder molecules, hydrophobic repulsion between drug and lubricant, and the mechanism of disintegrant water uptake and swelling.<sup>[17]</sup>

Students then enter a virtual GMP manufacturing environment and operate virtual wet granulation equipment. The simulation visualises granule formation in real time: drug and excipient powder particles are shown approaching each other, binder solution bridges forming between particles, and granules growing through coalescence and layering. The effect of granulation endpoint on granule size distribution, flowability, and compressibility is demonstrated by allowing students to vary binder concentration and granulation time and observing the consequences for granule quality in real time.<sup>[16]</sup>

Tablet compression is simulated with a virtual rotary tablet press in which students observe die filling, pre-compression, main compression, and ejection. The simulation visualises the effect of compression force on tablet porosity – showing the reduction in inter-particulate void space as compression force increases – and demonstrates how inadequate compression force produces friable tablets while excessive force causes capping and lamination. Tablet coating simulations show drug pellets entering the coating pan, rotating, and receiving spray coating solution that forms a polymer film; the simulation visualises the coalescence of spray droplets into a continuous film at the microscopic level.<sup>[16,17]</sup>

#### b) Drug Release and Dissolution Visualisation

One of the most educationally powerful applications of VR in pharmaceuticals is the visualisation of drug release mechanisms in controlled-release dosage forms – processes that are entirely invisible in a conventional dissolution apparatus. For a matrix tablet formulation, VR simulates the hydration and swelling of a hydroxypropyl methylcellulose (HPMC) matrix as dissolution medium penetrates the tablet surface. Students watch water molecules diffusing into the polymer matrix, the gradual gelatinisation of HPMC chains as they hydrate, and drug molecules diffusing through the increasingly viscous gel layer to reach the dissolution medium. The tortuosity of the diffusion pathway – the meandering route drug molecules must take through the gel layer – is visualised directly, making Higuchi's square root equation intuitive rather than abstract.<sup>[13,17]</sup>

For enteric-coated tablets, the simulation shows the tablet remaining intact in a virtual gastric environment (pH 1.2), with the enteric polymer coating resisting dissolution. As the tablet moves into a simulated intestinal environment (pH 6.8), students watch the ionisation state of the enteric polymer change, the coating dissolving from the outer surface inward, and drug release beginning only when the coating has fully dissolved. This direct visualisation of pH-dependent drug release mechanisms makes the rationale for enteric coating – protecting acid-labile drugs or delivering drugs to the intestine – immediately intuitive.<sup>[13]</sup>

#### c) Biopharmaceutics and Bioavailability Simulation

VR enables the simulation of the complete journey of a drug from oral administration to systemic circulation, visualising each step of the LADME sequence at mechanistic depth. Students watch drug particles in a virtual gastrointestinal tract dissolving according to the Noyes-Whitney equation – observing how particle size reduction increases dissolution rate by increasing surface area. Drug molecules in solution are shown partitioning across the intestinal epithelium through passive transcellular diffusion, active transport via membrane transporters such as PEPT1, and efflux by P-glycoprotein. The simulation visualises P-gp actively transporting drug molecules back into the intestinal lumen, demonstrating its role as a barrier to bioavailability in a way that textbook diagrams cannot.<sup>[13,17]</sup>

First-pass hepatic metabolism is simulated by showing drug molecules entering portal blood, reaching hepatocytes, and encountering cytochrome P450 enzymes in the endoplasmic reticulum. The simulation visualises enzyme-drug binding, metabolic transformation, and the production of metabolites, with the fraction escaping metabolism represented as the hepatic extraction ratio. The resulting plasma concentration-time profile is generated in real time, with students able to manipulate bioavailability-determining parameters – particle size, dissolution rate, food effect, enzyme induction – and immediately observe the consequences for C<sub>max</sub>, T<sub>max</sub>, and AUC.<sup>[13]</sup>

#### d) Quality Control and GMP Simulation

VR replicates complete quality control workflows for pharmaceutical products. Students perform virtual dissolution tests, observing tablets disintegrating and drug dissolving in the vessel while the dissolution apparatus records UV absorbance in real time to generate a dissolution profile. Deliberately defective batches – tablets with insufficient disintegrant, products with coating defects, parenteral preparations with visible particles – are presented to students who must identify the defect through virtual testing and root-cause analysis. Clean room behaviour simulations teach students GMP gowning procedures, airflow management, and contamination control protocols through immersive walkthroughs of ISO-classified environments.<sup>[16]</sup>



Fig 3: VR in Pharmaceuticals.

### 3 Advantages of VR in Pharmaceutics

VR provides access to pharmaceutical manufacturing environments that most academic institutions cannot physically replicate, dramatically reducing the gap between academic training and industry expectations. The ability to visualise drug release mechanisms at the molecular level – observing drug diffusion through polymer matrices, dissolution at particle surfaces, and membrane transport – provides mechanistic understanding that directly supports rational formulation design, a cognitive skill that cannot be developed from macroscopic laboratory observation alone. The consequence-free virtual environment also allows students to deliberately make formulation errors and observe their consequences without wasting materials or producing unusable products.<sup>[16,17]</sup>

### 4 Limitations of VR in Pharmaceutics

Pharmaceutics training requires development of genuine tactile skills – assessing granule texture, judging tablet hardness, detecting pour point anomalies in semi-solid preparations – that cannot be acquired through VR simulation. Virtual manufacturing environments simplify the variability and unpredictability of real manufacturing processes; real pharmaceutical manufacturing involves troubleshooting unexpected equipment behaviour, managing environmental variables, and making real-time judgments that require physical presence and sensorimotor experience. Platform-specific equipment simulations may not match the specific machines students encounter in industry, limiting transfer of learning.<sup>[7,9]</sup>

## III: Pharmaceutical Chemistry

### 1 Introduction to VR in Pharmaceutical Chemistry

Pharmaceutical Chemistry spans organic chemistry, medicinal chemistry, and analytical chemistry as applied to the discovery, synthesis, and quality control of drug substances. Practical training in this stream involves organic synthesis, identification and characterisation of drug molecules, and quantitative and qualitative analysis using chromatographic and spectroscopic techniques. The laboratory environment of pharmaceutical chemistry is characterised by hazardous chemicals, reactive reagents, flammable solvents, and toxic intermediates. For inexperienced students, the combination of chemical hazards and technically demanding procedures creates a genuine risk of accidents, making pharmaceutical chemistry one of the streams where VR's safety advantage is most compelling.<sup>[18]</sup>

Beyond safety, VR offers pharmaceutical chemistry education something genuinely revolutionary: the ability to observe chemical reactions at the molecular and electronic level. A student watching a nucleophilic substitution reaction in a conventional laboratory sees only a colour change and possibly a precipitate. In a VR pharmaceutical chemistry simulation, the same student watches electron density shift from the nucleophile toward the electrophilic carbon, visualises the HOMO-

LUMO orbital interaction that drives the reaction, observes the inversion of configuration at the stereogenic centre (Walden inversion), and tracks the formation and collapse of the pentacoordinate transition state. This is a transformation in the depth at which pharmaceutical chemistry can be taught and understood.<sup>[20]</sup>

## 2 Specific VR Applications in Pharmaceutical Chemistry

### a) Molecular-Level Reaction Visualisation

VR pharmaceutical chemistry platforms render organic reaction mechanisms at atomic and electronic resolution. Students can observe the SN2 reaction involved in the synthesis of a drug intermediate by watching bond polarisation, nucleophile approach along the correct trajectory relative to the leaving group (180° back-side attack), the simultaneous formation of the new bond and breaking of the old bond, and the umbrella inversion of the carbon configuration. Electron density is represented through colour-coded molecular electrostatic potential (MEP) surfaces that show how charge distribution changes dynamically during the reaction – the incoming nucleophile appearing red (electron-rich) and the electrophilic carbon appearing blue (electron-poor).<sup>[20]</sup>

For aromatic reactions such as electrophilic aromatic substitution – fundamental to the synthesis of many sulfonamide and NSAID drugs – students watch the pi electron cloud of the aromatic ring acting as a nucleophile toward the electrophile, the formation of the sigma complex (arenium ion) with loss of aromaticity, and the restoration of aromaticity through proton loss. The simulation visualises the resonance structures of the arenium ion dynamically, showing positive charge delocalisation across ortho and para positions – making the directing effects of ring substituents intuitively comprehensible rather than empirically memorized.<sup>[18,20]</sup>

### b) Drug Synthesis Procedures in VR

Students can perform step-by-step virtual synthesis of pharmacologically important molecules. The synthesis of aspirin (acetylsalicylic acid) from salicylic acid and acetic anhydride is simulated with full procedure fidelity: students add virtual reagents to a round-bottom flask, attach a virtual reflux condenser, heat the mixture to the correct temperature, observe the acylation reaction mechanism at the molecular level, perform virtual recrystallisation including hotfiltration and slow cooling crystallisation, and collect the virtual product. The simulation provides yield calculations and purity assessments based on the student's procedural choices, with deviations from optimal technique resulting in lower yield or impure product.<sup>[18]</sup>

The synthesis of sulfonamides, local anaesthetics such as benzocaine, and antipyretics such as paracetamol can all be simulated with equivalent procedural and mechanistic detail. For multi-step syntheses, VR platforms can track impurities through each synthetic step, demonstrating how an error in one step propagates through subsequent

steps to affect final product quality – a systems-level understanding of synthesis quality that is difficult to convey in a conventional laboratory setting.<sup>[19]</sup>

### c) Spectroscopic Analysis and Instrument Simulation

VR simulates the complete operation of analytical instruments used in pharmaceutical chemistry quality control. Students operate a virtual HPLC system: selecting a column, preparing mobile phase, injecting a sample, and interpreting the resulting chromatogram to identify and quantify drug content. The simulation visualises separation at the molecular level – showing drug molecules and impurities partitioning between the stationary phase and mobile phase in the column based on their polarity – making the principle of chromatographic separation tangible rather than abstract.<sup>[19]</sup>

Infrared spectroscopy is simulated by allowing students to 'record' a virtual IR spectrum of a drug molecule, then manipulate the three-dimensional molecular model to observe which functional groups are vibrating at the frequencies corresponding to absorption peaks. Students can click on an absorption peak in the spectrum and the simulation highlights the corresponding bond in the molecular model and animates its stretching or bending vibration – making the interpretation of IR spectra a genuinely three-dimensional and mechanistic exercise. UV-Vis spectroscopy, NMR spectroscopy, and mass spectrometry simulations follow similar principles.<sup>[19,20]</sup>

### d) Structure-Activity Relationship (SAR) and Medicinal Chemistry

VR medicinal chemistry platforms allow students to perform virtual drug design exercises by systematically modifying drug molecular structures and observing the consequences for receptor binding affinity and selectivity. Students select a lead compound, view its three-dimensional binding pose in the receptor active site, and then make specific structural modifications – adding a methyl group, replacing a hydroxyl with a fluorine, extending an alkyl chain – and immediately observe the computed change in binding energy and binding pose.<sup>[20]</sup>

This interactive SAR exploration makes the principles of bioisosterism, lipophilicity optimisation, and pharmacophore identification directly experiential. The molecular hydrophobicity surface of the drug is visualised in the context of the receptor binding pocket, making it immediately apparent why a modification that increases lipophilicity improves binding to a hydrophobic pocket. Platforms such as Molecular Workbench and Virtual Chem Lab implement these capabilities for multiple drug-receptor systems including beta-lactam antibiotics and their transpeptidase targets.<sup>[20]</sup>



Fig 4: VR in Pharmaceutical Chemistry.

### 3 Advantages of VR in Pharmaceutical Chemistry

The transformative advantage of VR in pharmaceutical chemistry is the ability to observe chemical processes at the molecular and electronic level – the dimension of reality where chemistry actually occurs. Conventional chemistry education requires students to infer molecular-level events from macroscopic observations; VR eliminates this inferential step by making molecular events directly observable. Multiple studies have documented improved understanding of reaction mechanisms and structural chemistry following VR-based molecular visualization.<sup>[20]</sup> Chemical safety is the other paramount advantage: students can learn to handle reactive and hazardous reagents, practice titration endpoint detection, and perform reflux procedures in a consequence-free virtual environment before encountering real chemical hazards.<sup>[18,19]</sup>

### 4 Limitations of VR in Pharmaceutical Chemistry

Pharmaceutical chemistry is ultimately a manipulative discipline requiring genuine dexterity with glassware, precision in weighing and volumetric operations, and judgment in assessing reaction progress by sensory observation. These manual laboratory skills cannot be developed through VR simulation. Virtual chemistry simulations also necessarily simplify real chemical complexity: real reactions produce by-products, real reagents have variable purity, and real chromatographic separations require troubleshooting that VR simulations do not replicate. The computational models underlying VR chemistry simulations are approximations of quantummechanical reality, and students must understand the distinction between VR representations and actual molecular structure.<sup>[18,19]</sup>

## IV: Pharmaceutical Biotechnology

### 1 Introduction to VR in Pharmaceutical Biotechnology

Pharmaceutical Biotechnology applies biological principles and molecular biology techniques to the development, production, and quality control of biopharmaceuticals – drugs derived from living organisms including recombinant proteins, monoclonal antibodies, vaccines, and gene therapy vectors. This stream is among the most rapidly advancing in pharmaceutical sciences and is increasingly central to drug development globally. Practical training in

biotechnology requires cell culture facilities, biosafety cabinets, fermenters, PCR thermocyclers, electrophoresis equipment, ELISA readers, Western blot systems, and a range of molecular biology instrumentation that collectively represents capital investment far beyond the capacity of most pharmacy colleges.<sup>[21]</sup>

The biosafety requirements of biotechnology education add another layer of complexity. Working with recombinant organisms, cell lines, and immunological materials requires biosafety level (BSL) 2 infrastructure, institutional biosafety committee approval, and trained personnel. VR simulation addresses both the infrastructure and biosafety barriers simultaneously, enabling students to perform detailed biotechnology procedures in an immersive virtual BSL-2 laboratory environment that would be prohibitively expensive and logistically complex to establish physically.<sup>[22]</sup>

## 2 Specific VR Applications in Pharmaceutical Biotechnology

### a) Genetic Engineering and Recombinant DNA Technology

The recombinant DNA technology workflow – the foundation of biopharmaceutical production – is one of the most powerful VR applications in pharmaceutical biotechnology. Students perform a complete virtual recombinant protein production experiment beginning with restriction enzyme digestion. In the VR simulation, students select a restriction enzyme such as EcoRI and add it to a virtual plasmid vector, then watch at the molecular level as the enzyme scans the double-stranded DNA sequence, recognises its palindromic recognition sequence (5'-GAATTC-3'), and makes staggered cuts in both strands to produce cohesive ('sticky') ends. The three-dimensional structure of the restriction enzyme bound to DNA is visualised, making the structural basis of sequence-specific recognition directly observable.<sup>[22]</sup>

The ligation step is simulated with equivalent molecular detail: students watch the complementary cohesive ends of the digested plasmid and insert DNA annealing through hydrogen bond formation between complementary base pairs, then observe DNA ligase catalysing the formation of the phosphodiester bond that seals the nicks in the DNA backbone, producing a covalently closed recombinant plasmid. Transformation of competent *Escherichia coli* cells is simulated, showing the competent cell in a calcium chloride solution, the heat shock step causing transient pore formation in the cell membrane, and the recombinant plasmid entering the cell and establishing within the cytoplasm.<sup>[22,24]</sup>

Colony selection, plasmid isolation, and confirmation of recombinant insert by restriction digestion and gel electrophoresis are all simulated with procedural detail. The virtual gel electrophoresis simulation shows DNA migrating through the agarose matrix under the influence of the electric field, with smaller fragments moving faster through the gel pores –making the physical basis

of size-dependent separation intuitive through direct animated visualization.<sup>[22]</sup>

### b) PCR and Gene Amplification

The polymerase chain reaction (PCR) is simulated at the molecular level through all three thermal cycling stages. During the denaturation step (95°C), students watch the hydrogen bonds between DNA base pairs breaking and the double helix unwinding into two single-stranded templates. During the annealing step (55-65°C), forward and reverse primers are shown approaching their complementary sequences on the template strands, forming hydrogen bonds in an antiparallel orientation, and establishing the 3'-OH terminus required for polymerase extension. During the extension step (72°C), Taq DNA polymerase is visualised binding to the primer-template junction, incorporating deoxyribonucleotide triphosphates (dNTPs) from the virtual reaction mixture in the 5'-to-3' direction, and synthesising the new complementary strand.<sup>[23]</sup>

The exponential amplification of the target sequence across 30-35 cycles is animated, with the number of target copies doubling each cycle from 1 to 2 to 4 to 8, and so on to over a billion copies. Students can manipulate primer design, annealing temperature, and MgCl<sub>2</sub> concentration and observe the effects on PCR efficiency and specificity, developing the troubleshooting skills essential for research and quality control applications.<sup>[23]</sup>

### c) Cell Culture and Bioreactor Simulation

Cell culture simulations replicate the aseptic conditions and procedural discipline of real biological laboratories. Students enter a virtual BSL-2 laboratory, don virtual personal protective equipment following correct gowning sequence, and work within a virtual Class II biosafety cabinet. Subculturing of a Chinese Hamster Ovary (CHO) cell line used for recombinant protein production is simulated: students observe the monolayer of virtual cells under a virtual microscope, assess confluency, remove spent medium, wash with virtual PBS, add trypsin, and observe cell detachment. The simulation visualises trypsin cleaving cell surface adhesion proteins at the molecular level, explaining why trypsinisation disrupts cell monolayers.<sup>[21]</sup>

Bioreactor operation for large-scale protein production is simulated with full process parameter monitoring. Students observe the virtual fermenter vessel, set agitation speed, aeration rate, temperature, and pH, and monitor dissolved oxygen tension and cell density through virtual sensors. The simulation demonstrates how inadequate aeration leads to hypoxic conditions and cell death – visualising declining dissolved oxygen, increasing lactate accumulation, and cell necrosis. Fed-batch feeding strategies are simulated, with students determining optimal nutrient addition timing based on cell growth parameters.<sup>[21,22]</sup>

#### d) Immunology and Vaccine Technology

Immunological processes fundamental to vaccine development and monoclonal antibody production are visualised in VR with a mechanistic depth impossible in conventional laboratory instruction. Students explore a three-dimensional simulation of the lymph node during antigen presentation: dendritic cells are shown processing antigen, loading peptide fragments onto MHC class II molecules, and presenting them to T helper lymphocytes in the paracortex. The immunological synapse between APC and T cell is visualised at the molecular level, showing TCR-pMHC interaction and co-stimulatory signal engagement.<sup>[24]</sup>

The humoral immune response is animated sequentially: T cell activation, cytokine signalling to B cells, B cell differentiation into plasma cells, affinity maturation in germinal centres, and the production of high-affinity IgG antibodies. The mechanism of monoclonal antibody production by hybridoma technology is simulated – showing B cell fusion with myeloma cells, HAT selection medium eliminating unfused myeloma cells, and hybridoma clones secreting monoclonal antibody.<sup>[21,24]</sup>



Fig. 5: VR in Biotech.

### 3 Advantages of VR in Pharmaceutical Biotechnology

VR democratizes access to biotechnology laboratory environments that most academic institutions cannot physically replicate. Students at colleges without cell culture facilities or molecular biology equipment can gain detailed procedural familiarity with recombinant DNA technology, PCR, and cell culture through VR simulation. The elimination of biosafety risk makes it possible to visualise and practice procedures involving recombinant organisms and biological materials that would require BSL-2 containment in a real laboratory. The molecular-level visualisation of genetic engineering processes – restriction enzyme recognition, primer annealing, polymerase extension – transforms abstract biochemical concepts into directly observable phenomena.<sup>[22,23,24]</sup>

### 4 Limitations of VR in Pharmaceutical Biotechnology

Biotechnology laboratory competency ultimately depends on developing aseptic technique discipline – the constant vigilance, physical habits, and sensorimotor

automaticity required to work safely with biological materials without contamination. These skills cannot be developed through VR simulation alone. Real cell culture involves unpredictable biological variability: cells grow at variable rates, contamination events occur unexpectedly, and bioreactor processes develop in ways that require real-time troubleshooting. The interpretation of real experimental data – unexpected PCR bands, anomalous ELISA results, inconsistent bioreactor performance – requires exposure to real laboratory systems.<sup>[21,22]</sup>

### V: Biochemical Applications of VR in Pharmacy Education

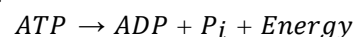
#### 1. Introduction to Biochemical-Level Visualisation

Biochemistry forms the fundamental basis of all physiological and pharmacological processes, involving complex molecular interactions such as enzyme catalysis, metabolic pathways, and energy transfer systems. Traditional teaching methods rely heavily on static diagrams and pathway charts, which often fail to convey the dynamic and interconnected nature of biochemical reactions. Virtual Reality (VR) provides a transformative platform for visualising these biochemical processes in real time, enabling students to observe molecular events at a depth that is not possible in conventional laboratory or classroom settings.

#### 2. Specific VR Applications in Biochemistry

##### a) Cellular Energy Metabolism (ATP–ADP Cycle)

VR allows students to visualise **cellular energy transfer mechanisms** in real time. Students can observe the hydrolysis of ATP into ADP and inorganic phosphate (Pi), with the release of energy used for cellular processes such as active transport and muscle contraction.



The simulation demonstrates:

- Breaking of high-energy phosphate bonds
  - Energy release and coupling with cellular work
  - Regeneration of ATP via oxidative phosphorylation
- Students can further explore mitochondrial processes such as:
- Electron transport chain (ETC)
  - Proton gradient formation
  - ATP synthase activity

##### b) Enzyme Kinetics and Drug–Enzyme Interaction

VR platforms simulate enzyme-substrate interactions using dynamic 3D models. Students can observe:

- Formation of enzyme-substrate complex
- Active site binding and specificity
- Competitive and non-competitive inhibition

$$v = \frac{V_{max} [S]}{K_m + [S]}$$

This helps in understanding:

- Mechanism of enzyme inhibition by drugs

- Role of enzymes in drug metabolism
- Pharmacokinetic implications

#### c) Metabolic Pathway Visualisation

VR enables immersive exploration of key metabolic pathways such as:

- Glycolysis
- Citric acid cycle (Krebs cycle)
- Lipid metabolism
- Amino acid metabolism

Students can follow each step of glucose metabolism from glucose to pyruvate and further into the TCA cycle, observing:

- Enzyme-mediated transformations
- Intermediate formation
- Energy yield (ATP, NADH, FADH<sub>2</sub>)

#### d) Signal Transduction Pathways

VR simulations provide detailed insights into intracellular signalling mechanisms such as:

- G-protein coupled receptor (GPCR) pathways
- Second messenger systems (cAMP, IP<sub>3</sub>, DAG)
- Protein phosphorylation cascades

Students can observe how extracellular signals lead to intracellular biochemical responses, linking pharmacology with biochemistry at a molecular level.

#### e) Drug Metabolism and Biotransformation

VR allows visualisation of hepatic drug metabolism processes including:

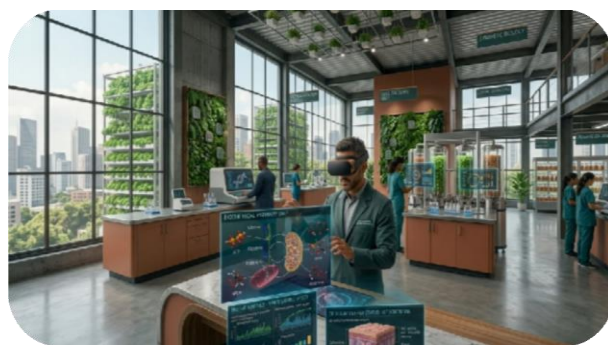
- Phase I reactions (oxidation, reduction, hydrolysis)
- Phase II reactions (conjugation) Students can observe:
- Cytochrome P450 enzyme activity
- Formation of metabolites
- Detoxification vs activation of prodrugs

### 3. Advantages of VR in Biochemical Applications

An important advantage that VR technology provides for biochemical learning is the potential to change the abstract, dynamic process of biochemical reactions to one which can actually be witnessed in real time. For example, in addition to learning about ATP hydrolysis, enzymes, and metabolic cycles via two-dimensional drawings and models, virtual reality allows students to see these chemical reactions taking place, with the interactions between bonds and the energy transfer that accompanies these processes visible at all times. Virtual reality helps to improve the student's understanding of the process involved in enzyme kinetics by allowing them to view these processes instead of merely inferring their results. VR also facilitates the connection between biochemical reactions and their effects in pharmacology and physiology, with drug reactions being traced from receptor interactions to their metabolic consequences in terms of enzyme inhibition and energy production.

### 4. Limitations

However, apart from the numerous advantages of VR-based biochemical simulation, it still fails to provide sufficient practice due to the high degree of complexity and interconnection of biological systems that are regulated by feedback mechanisms and affected by specific cellular conditions. In other words, biological processes like glycolysis and enzyme activity may be shown as linear sequences, making it less probable for users to be faced with biological diversity. Furthermore, VR can be used only to gain qualitative knowledge since it does not allow practicing skills that cannot be learned without practical experience, such as enzyme assays, spectrophotometry, chromatography, and kinetic data analysis. Currently, available software products do not cover all aspects of biochemical processes and drug metabolism; besides, the accuracy of results is provided by computations only. Finally, the lack of opportunities to face actual difficulties in a biochemical laboratory will result in insufficient experience in handling problems.



**Fig. 6: VR in Biochem.**

## VI: Clinical Pharmacy

### 1 Introduction to VR in Clinical Pharmacy

Clinical Pharmacy is the patient-centred practice stream that prepares pharmacy graduates for pharmaceutical care roles in hospital, community, and outpatient settings. Clinical pharmacy competencies include medication therapy management, drug information provision, patient counselling, adverse drug reaction monitoring, prescription review, drug interaction identification, and interprofessional healthcare team collaboration. Unlike laboratory-based pharmaceutical streams, clinical pharmacy training is fundamentally a communication, reasoning, and decision-making discipline in which students must learn to interact with diverse and often vulnerable patient populations in real clinical environments.<sup>[12]</sup>

The provision of authentic clinical training experiences to pharmacy students is inherently constrained by patient safety considerations, institutional access, and the limited supervision capacity of clinical preceptors. Real patients cannot be placed at risk for student learning purposes, and student-supervised clinical encounters are too few in number to develop competency across the full range of clinical pharmacy scenarios. VR clinical simulation addresses this gap by providing unlimited access to

realistic patient encounters without risk to real patients or to the student's professional record.<sup>[14,25]</sup>

## 2 Specific VR Applications in Clinical Pharmacy

### a) Patient Counselling Simulation

VR patient counselling simulations use artificially intelligent virtual patients that respond dynamically and realistically to the student's communication choices. The student, represented by an avatar in the virtual environment, approaches a virtual patient in a simulated pharmacy consultation room. The patient presents with a new prescription for a drug such as warfarin, and the student must conduct a structured medication counselling session – explaining the drug's purpose, correct administration, potential side effects, monitoring requirements (INR), dietary interactions (vitamin K-rich foods), and when to seek medical attention.<sup>[12]</sup>

The AI-driven virtual patient responds differently based on the quality of information the student provides. If the student omits the interaction between warfarin and common OTC medications such as aspirin, the virtual patient may subsequently ask about taking aspirin for a headache, requiring the student to recognise and correct the potential interaction in real time. If the student's explanation is unclear or uses excessive medical terminology, the virtual patient expresses confusion and asks for clarification, training the student to adapt their communication style to patient health literacy. Scenarios can represent patients with hearing impairment, non-native language speakers, and patients with health anxiety, developing communication adaptability across diverse populations.<sup>[12,14]</sup>

### b) Prescription Error Detection and Drug Safety

Virtual pharmacy dispensing environments present students with prescription orders containing deliberate clinical errors embedded within otherwise realistic prescription data. Students must process each prescription through a complete verification workflow: confirming patient identity, reviewing prescribed drug, dose, route, frequency, duration, and indication; cross-checking against the patient's virtual medication history for drug interactions and duplications; screening for contraindications against documented allergies and comorbidities; and calculating dose correctness for weight-based and renal dose-adjusted medications.<sup>[14]</sup>

Examples of embedded errors include: a prescription for methotrexate specifying a daily dose rather than the correct once-weekly dose (a potentially fatal prescribing error in clinical practice); a paediatric prescription with an incorrect weight-based dose calculation; an anticoagulant prescribed to a patient whose virtual record shows a documented allergy to heparin; or a drug prescribed to a patient with a renal impairment requiring dose reduction. The VR system records the student's error-detection performance across a battery of such prescriptions, providing detailed feedback on error types missed and the clinical consequences that would have

resulted.<sup>[14,25]</sup>

### c) Clinical Ward and Hospital Pharmacy Simulation

Immersive hospital pharmacy VR environments replicate the workflow of a clinical pharmacy practitioner conducting ward rounds, medication reconciliation, and interprofessional team meetings. Students navigate a virtual ward, access virtual electronic patient records, review medication charts, and identify drug-related problems: potentially inappropriate medications in elderly patients per Beers Criteria, antimicrobial stewardship issues in patients receiving prolonged broad-spectrum antibiotics, pain management optimisation in post-operative patients, and therapeutic drug monitoring recommendations for narrow therapeutic index drugs such as vancomycin and aminoglycosides.<sup>[25]</sup>

Multiplayer VR environments simulate the interprofessional dynamics of real hospital practice. A virtual physician avatar may challenge the student's drug therapy recommendation, requiring the student to articulate the evidence base for their suggestion and negotiate a therapeutic decision. Virtual nursing avatars may report medication administration errors, requiring the student to manage the clinical consequences and initiate appropriate reporting procedures.<sup>[14]</sup>



Fig. 7: VR in Clinical Pharmacy.

## 3 Advantages of VR in Clinical Pharmacy

VR provides clinical pharmacy students with exposure to the full spectrum of clinical scenarios – including rare drug interactions, complex polypharmacy presentations, and serious prescribing errors – that cannot be systematically encountered in real clinical placements of equivalent duration. The AI-driven virtual patient enables personalised learning: the system can identify the specific communication skills or clinical reasoning gaps of each student and generate scenarios specifically designed to address those gaps. The elimination of patient risk removes the anxiety that frequently impairs student performance in real clinical encounters, producing better and more confident learners.<sup>[12,25]</sup> Multiple studies have documented improved prescription error detection rates, stronger patient counselling competency, and greater clinical confidence following VR-based clinical pharmacy training.<sup>[14]</sup>

## 4 Limitations of VR in Clinical Pharmacy

The non-verbal dimensions of real patient

communication – reading subtle facial cues, managing patient distress, building genuine therapeutic rapport – require real human interaction for adequate development and cannot be fully replicated by virtual patient AI. Real patients bring unique combinations of cultural background, emotional state, health beliefs, and health literacy that current virtual patient technology cannot represent with sufficient diversity and authenticity. Clinical pharmacy also requires exposure to real health record systems, institutional drug policy frameworks, and formulary management that vary between healthcare settings and cannot be generalised in VR simulations.<sup>[12,14]</sup>

## DISCUSSION

The stream-wise examination of VR applications across five pharmaceutical disciplines reveals a consistent pattern of educational benefit alongside discipline-specific opportunities and limitations. Across all five streams, VR technology demonstrates significant capacity to address the practical training challenges that characterise contemporary pharmacy education in India. The pattern that emerges is one of complementarity rather than replacement: in each stream, VR is most educationally powerful when it provides what the real laboratory cannot – molecular-level visualisation in pharmacology and chemistry, equipment access in pharmaceuticals and biotechnology, and patient exposure breadth in clinical pharmacy – while the real laboratory or clinical environment continues to develop the sensorimotor and interpersonal competencies that VR cannot simulate.<sup>[1,2,5]</sup>

The most compelling and specific contribution of VR, identified consistently across the pharmacology, chemistry, and biotechnology streams, is the ability to make invisible processes visible. The pharmacological mechanism of action of a drug is invisible in a conventional laboratory; VR makes it directly observable. The mechanism of a nucleophilic substitution reaction is invisible to a chemistry student watching a colour change; VR shows the electron movement, transition state geometry, and configuration inversion. The molecular specificity of restriction enzyme recognition is invisible during a gel electrophoresis experiment; VR shows the enzyme reading the DNA sequence. This transformation – from inferring invisible mechanisms to directly observing them – represents a qualitative improvement in the depth at which pharmaceutical science can be taught.<sup>[3,11,20,22]</sup>

The Indian pharmacy education context adds particular urgency to the case for VR adoption across all five streams. Regulatory tightening of animal use requirements creates acute need in pharmacology; capital cost barriers restrict pharmaceuticals and biotechnology equipment access; chemical safety concerns in pharmaceutical chemistry argue for pre-lab VR familiarisation; and clinical placement capacity limitations constrain clinical pharmacy training. VR

platforms accessible via mobile devices and low-cost HMD hardware can meaningfully address all five of these barriers simultaneously, and are particularly valuable for the substantial proportion of Indian pharmacy colleges that operate with limited infrastructure.<sup>[4,9,10]</sup>

The reference evidence reviewed across all five streams also highlights a consistent and significant gap: the near-total absence of VR platforms designed specifically for the Indian pharmacy curriculum. The B.Pharm and M.Pharm curricula approved by the Pharmacy Council of India specify particular experiments, drug classes, and practical competencies that differ from the Western curricula for which most commercially available VR pharmacy platforms were developed. A suite of India-specific VR pharmacy simulation tools – covering the specific experiments specified in the PCI curriculum, using drug molecules and experimental protocols relevant to Indian pharmacological practice, and delivered in a format compatible with the infrastructure available at Indian pharmacy colleges – would represent a significant and educationally transformative development.<sup>[4,5]</sup>

Future directions for VR in pharmacy education include AI-integrated adaptive learning systems that monitor individual student performance in each stream and dynamically adjust the virtual learning experience to address specific conceptual or procedural gaps; collaborative multiplayer VR environments that replicate the social dimensions of real laboratory and clinical practice; and extended reality platforms that blend VR's full immersion with AR's real-world contextual overlay in a unified experience. The formal integration of VR-based practical assessment into the Pharmacy Council of India's competency standards – establishing which practical competencies can be validly assessed through VR performance and which require real laboratory or clinical demonstration – is the next critical step toward mainstream VR adoption in Indian pharmacy education.<sup>[13,14,25]</sup>

## CONCLUSION

Virtual Reality technology represents a significant and evidence-supported opportunity to transform the quality, ethics, and accessibility of pharmacy practical education across all major pharmaceutical streams. The stream-wise analysis presented in this review identifies a consistent pattern: VR makes processes visible that are invisible in real laboratories, provides access to environments and equipment that most institutions cannot afford, and creates safe practice opportunities for procedures that are hazardous, irreversible, or ethically restricted in real settings.

In pharmacology, VR reveals the molecular pharmacodynamics that underlie observed physiological effects. In pharmaceuticals, it visualises drug dissolution, release kinetics, and bioavailability mechanisms at the molecular scale. In pharmaceutical chemistry, it makes reaction mechanisms and molecular interactions directly

observable at the electronic level. In pharmaceutical biotechnology, it enables immersive genetic engineering, cell culture, and immunology practicals in simulated BSL-2 environments. In clinical pharmacy, it provides unlimited access to realistic patient counselling, prescription error detection, and ward pharmacy practice scenarios.

Across all five streams, VR is most powerful as a complement to real practical experience –providing conceptual preparation, procedure familiarisation, and molecular-level insight that maximises the educational value of limited real laboratory and clinical exposure time. The development of India-specific VR platforms tailored to the Pharmacy Council of India curriculum, combined with formal regulatory recognition of VR-based competency assessment, would enable the documented educational benefits of immersive simulation to be systematically realised across the full spectrum of Indian pharmacy institutions, from well-resourced urban colleges to under-equipped rural institutions.

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#### REFERENCES

- Smith A, Brown T. Applications of virtual reality in medical education: a systematic review. *J Med Simul.*, 2018; 12(2): 85–92.
- Johnson P, Lee M. Immersive learning technologies in healthcare education: current applications and future directions. *Med Educ Rev.*, 2019; 23(4): 210–218.
- Kumar R, Sharma S, Patel K. Virtual laboratory simulation in pharmacology education: a review of platforms and learning outcomes. *Int J Pharm Educ Res.*, 2020; 11(3): 145–150.
- Patel N, Desai K, Shah P. Technology-assisted learning tools in Indian pharmacy education: challenges and opportunities. *J Pharm Sci Res.*, 2021; 13(5): 289–295.
- Sharma V, Gupta R, Singh A. Virtual simulation models for biomedical and pharmaceutical training: compliance with 3Rs and CPCSEA guidelines. *Int J Pharm Sci Rev Res.*, 2022; 74(1): 102–107.
- Radianti J, Majchrzak TA, Fromm J, Wohlgenannt I. A systematic review of immersive virtual reality applications for higher education: design elements, lessons learned, and research gaps. *Educ Inf Technol.*, 2020; 25(2): 1535–1571.
- Moro C, Stromberga Z, Raikos A, Stirling A. The effectiveness of virtual and augmented reality in health science education: a systematic review. *Anat Sci Educ.*, 2017; 10(6): 549–559.
- Makransky G, Petersen GB. Immersive virtual reality increases liking but not learning. *Learn Instruct.*, 2019; 31(4): 895–916.
- Kyaw BM, Saxena N, Posadzki P, et al. Virtual reality for health professions education: systematic review and meta-analysis by the Digital Health Education Collaboration. *J Med Internet Res.*, 2019; 21(1): e12959.
- Pantelidis VS. Reasons to use virtual reality in education and training courses and a model to determine when to use virtual reality. *Themes Sci Technol Educ.*, 2010; 2(1–2): 59–70.
- Merchant Z, Goetz ET, Cifuentes L, Keeney-Kennicutt W, Davis TJ. Effectiveness of virtual reality-based instruction on students' learning outcomes in K-12 and higher education: a meta-analysis. *Comput Educ.*, 2014; 70: 29–40.
- Jensen L, Konradsen F. A review of the use of virtual reality head-mounted displays in education and training: pitfalls and possibilities. *Educ Inf Technol.*, 2018; 23(4): 1515–1529.
- Freina L, Ott M. A literature review on immersive virtual reality in education: state of the art and perspectives. In: *Proceedings of eLearning and Software for Education Conference*, 2015; 1: 133–141.
- Makransky G, Petersen GB. The cognitive affective model of immersive learning (CAMIL): a theoretical research-based model of learning in immersive virtual reality. *Educ Psychol Rev.*, 2021; 33(3): 1105–1134.
- Parong J, Mayer RE. Learning science in immersive virtual reality. *J Educ Psychol.*, 2018; 110(6): 785–797.
- Lam K, Madhavan S, Sunder S. Virtual reality simulations in pharmaceutical manufacturing education: a scoping review. *J Pharm Technol.*, 2020; 36(4): 178–186.
- Fitzgerald E, Lawton J, Buss A. Using virtual simulation to teach pharmaceutical sciences: student perceptions and learning outcomes. *Curr Pharm Teach Learn*, 2021; 13(7): 856–863.
- Winkelmann K, Scott MF, Wong DL. A study of high school students' performance of a chemistry experiment within a virtual laboratory system. *J Chem Educ.*, 2014; 91(9): 1336–1340.
- Roach LO, Scherz JW. Virtual reality for teaching organic chemistry: impact on mechanistic understanding, spatial reasoning and student attitude. *J Chem Educ.*, 2019; 96(3): 453–461.
- Dori YJ, Barak M. Virtual and physical molecular modelling: fostering model perception and spatial understanding in chemistry and pharmaceutical sciences. *Educ Technol Soc.*, 2001; 4(1): 61–74.
- Grantcharov TP, Bardram L, Funch-Jensen P, Rosenberg J. Learning curves and impact of previous operative experience on performance on a virtual reality simulator. *Am J Surg.*, 2003; 185(2):

- 146–149.
22. Labster ApS. Virtual laboratory simulations for biotechnology and pharmaceutical sciences: platform overview and impact data. Labster Impact Report. Copenhagen: Labster, 2022.
  23. Bransford JD, Brown AL, Cocking RR, editors. How People Learn: Brain, Mind, Experience, and School. Expanded ed. Washington DC: National Academy Press, 2000.
  24. Dede C. Immersive interfaces for engagement and learning in the biological and pharmaceutical sciences. *Science*, 2009; 323(5910): 66–69.
  25. Clauson KA, Aungst TD, Sherrill B. Clinical pharmacy education in the digital age: evaluating the potential role of virtual reality for patient counselling and medication management training. *Am J Pharm Educ.*, 2019; 83(4): 6857.